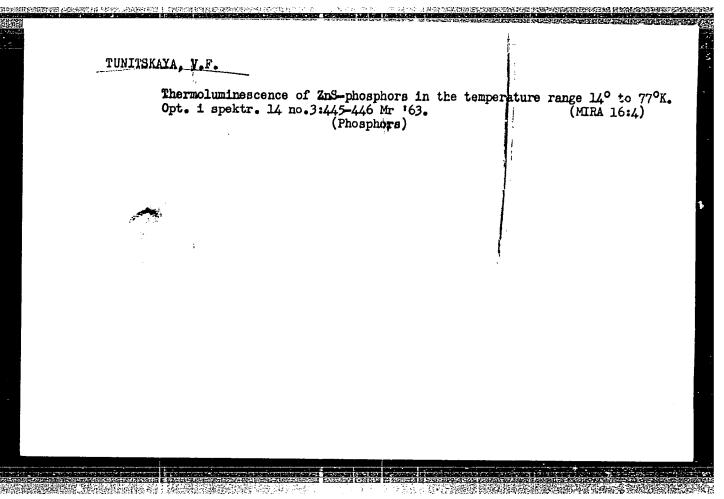
L 19486-63 EWT(1)/EWP(a)/EWT(E)/EWP(B)/BDS _AFTTC/ASD/IJP(C)/SSD S/2911/63/001/000/0290/0299 ACCESSION NR: AT3002237 AUTHORS: Levshin, V. L.; Reshetina, T. S.; Tunitskaya, V. F.; Vesil'yeva. TITLE: Stimulating action of infrared radiation on zinc sulfide phosphors SOURCE: Optika i spektroskopiya; sbornik statey. v. 1: Lyuminestsentsiya. Moscow. Izd-vo AN SSSR, 1963, 290-299 TOPIC TAGS: electron, tr p, energy level, infrared, absorption, flashing, phosphor /escence ABSTRACT: An investigation was made of the flashing process in ZnS with electrons trapped (or localized) in shallow levels under infrared excitation of wavelength 1 µ to 3.5 \(\mu \). The infrared response of these phosphors was studied at -77, -196 and -259C. Flash-emitting energy levels were established after obtaining the thermoluminescence curves of several zinc sulfide phosphors. The effect of infrared radiation of various wave lengths on one specimen, under varying conditions of excitation, was studied in greet detail. It is shown that quenching, maximum absorption in radiation spectra, and the flesh magnitude under stimulation of infrared radiation at the excitation; level of 365 millimicron is 1.5 to 2.0 times lower than the excitation at λ = 312 millimicron. This is attributed to action of p-type levels Card 1/2

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S/051/63/014/003/019/019 E039/E120

AUTHOR:

Tunitskaya, V.P.

TITLE:

Thermal luminescence of ZnS - phosphors in the

temperature range 14 - 77 K

PERIODICAL: Optika i spektroskopiya, v.14, no.3, 1963, 445-446

TEXT: A special helium cryostat of the type described by E.N. Lotkova and A.B. Fradkov (PTE no.1, 1961, 188) is used in these experiments. The ZnS samples are in the form of a 4 µ thick layer on duralumin (1.5 mg/cm) mounted on a resistance the mometer which is tightly screwed to a cold conductor. A heating coil is situated between the sample and the helium bath, and increases the sample temperature at a rate of 9 min. Luminescence is detected on a photomultiplier and the 365 mµ line is used for excitation. Excitation time at 14 k is 10 minutes; the sample is then heated after a period of one minute from the time of completion of excitation. ZnS phosphors of various types are investigated. It is shown that the shape of the thermal luminescence curve is independent of the presence of copper in the range 14 - 80 k.

Card 1/2

Thermal luminescence of ZnS - ... \$/051/63/014/003/019/019 E039/E120

and shows a weak peak at about 70 °K. Similarly the curve for ZnS - Cu heated in molten MgCl₂, BaCl₂ and ZnCl₂ shows the same typical shape. ZnS - Ag heated in an atmosphere of HCl produced a broader peak with a maximum at about 60 °K. These results show that the ZnS phosphors have a small trapping level. Samples with a hexagonal modification have a background extending down to 14 °K. This effect will be the subject of a further paper. At the present time it is assumed that the small levels are not connected with any activator, but depend only on the structure of the basic material. There is 1 figure.

SUBMITTED: October 20, 1962

Card 2/2

LEVSHIN, V.L.; TUNITSKAYA, V.F.

Luminescence of ZnS-Mn phosphors during excitation and its kinetics.

Opt. 1 spektr. 9 no.2:223-232 Ag '60. (MIRA 13:8)

(Luminescence) (Phosphors)

LEYSHIN, V.L.; TUNITSKAYA, V.F.

Thermoluminescence and localization levels of ZnS-Mn phosphore.
Opt. i spektr. 8 no.5:663-671 My '60. (MIRA 13:9)
(Zinc sulfide) (Luminescence)

s/051/60/008/005/011/027 E201/E491

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AUTHORS 8

Levshin, V.L. and Tunitskaya, V.F. Thermal De-excitation and Localization Levels of

TITLE:

PERIODICAL: Optika i spektroskopiya, 1960, Vol.8, No.5, pp.663-671

A series of ZnS-Mn phosphors was studied in a wide range of temperatures using exciting light of various wavelengths and The first part of this investigation is reported in the present paper which deals with the structure, population and thermal de-excitation (emptying) of capture levels in ZnS-Mn The effect of increasing manganese concentration on the spectrum of the levels was studied, as well as the effect of the excitation wavelengths on the population of the levels. The phosphors were prepared by heating a charge in a stream of nitrogen at 850°C for 30 min. Manganese was introduced Stream of Mich and its initial concentration in the charge in the form of Mich and its initial concentration in the charge was from 10^{-6} to 3×10^{-2} g/g. About 40% of Min was lost during X-ray diffraction studies showed that all samples had Some of the results are presented in firing. sphalerite structure.

Card 1/4

S/051/60/008/005/011/027 E201/B491

Thermal De-excitation and Localization Levels of ZnS-Mn Phosphors

Fig.1 and 2 show the thermoluminescence (thermal de-excitation) curves for phosphors prepared at various temperatures (Fig.1) and with various amounts of manganese (Fig. 2). Fig. 3 and 4 and Tables 1 and 2 give the analysis of peaks I, II and III in thermoluminescence curves. Fig. 5 and 6 show the light sums of blue, manganese and total luminescence as a function of the concentration of manganese. Fig.7 shows the thermoluminescence curves of ZnS (Fig.7a) and of ZnS with 10-3 g/g Mn (Fig.7b) excited with light of 312 (curves 1). luminescence light sums when the phosphors were excited with light 366 (curves 2) and 436 mm (curves 3). of 436 mm wavelength (compared with excitation using 312 and 366 mm drawn from the results: (1) Manganese itself does not produce new localization levels which would appear in thermoluminescence peaks Only a small number of Mn ions remains near places where electrons are localized by oxygen and these Mn ions deepen the oxygen localization levels producing a new peak at -30°C. Card 2/4

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S/051/60/008/005/011/027 E201/E491

Thermal De-excitation and Localization Levels of ZnS-Mn Phosphors

Formation of this new peak is accompanied by a reduction of the oxygen peak at -90°C. (2) The quantum light sums of ZnS-Mn with up to 6×10^{-3} g/g Mn are practically independent of the concentration of Mn. In such phosphors radiationless transitions of electrons from localization levels do not occur on thermal de-excitation. (3) Light of 312 and 366 mm wavelengths does not affect localized electrons and produces similar light sums on thermal de-excitation. If light of $\lambda = 436 \text{ mm}$ is used, light sums are reduced by factor of 3 and this reduction is stronger in the case of the blue luminescence compared than for orange luminescence; luminescence due to shallow levels is weakened to a greater extent than luminescence due to deep levels. (4) Increase of the amount of Mn reduces the depth of electron localization levels. (5) The prcfile of the high-temperature thermoluminescence peak suggests that only infrequently electrons are recaptured by localization levels during the process of thermoluminescence. Acknowledgments are made to A.V. Lavrov for preparation and analysis of the phosphors and to O.M. Agafonova for Card 3/4

S/051/60/008/005/011/027
Thermal De-excitation and Localization Levels of ZnS-Mn Phosphors
her help in experiments.
13 references; 11 Soviet, 1 German and 1 Dutch.

SUBMITTED: July 25, 1959

Card 4/4

Card 1/2

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S/051/60/009/002/002/006 E201/E691

AUTHORS:

Levshin, V.L. and Tunitskaya, V.F.

TITLE:

The Luminescence Processes and their Kinetics in ZrS-Mn Phosphore During Excitation

PERIODICAL: Optika i spektroskopiya, 1960, Vol. 9, No. 2, pp. 223-232

TEXT: In an earlier paper (Ref 1) the authors dealt with thermal deexcitation and the origin of localization levels in ZnS-Mn phosphors. The present paper extends this work to luminescence of the same phosphors under excitation. The phosphors were in the form of $\sim 3.7~\mu$ thick layers (with grain diameter of $\sim 1~\mu$) and were excited with 365 and 312 mm lines from a mercury lamp; the quantum intensities of the lines were equalized with filters. The intensity of luminescence was measured with a photenultiplier FEU-32. The author studied the effect of temperature on the absorption spectra (Figs. 1, 2) and on the blue and orange luminescence of the phosphors (Figs. 3-6). The information obtained is used to discuss the processes of transformation of the excitation energy into blue luminescence, manganese luminescence and heat, as a function of the ambient temperature and the concentration of manganese.

Card 1/2

S/051/60/009/002/002/005 B201/E691

The Luminescence Processes and their Kinetics in ZnS-Mn Phosphors During Excitation

The mechanism of luminescence and quenching processes are dealt with and an energy diagram is proposed (Fig. 7) for the localization levels of electrons and holes. Acknowledgments are made to 0.M. Agafonova for her help in measurements and to A.V. Lavrov for preparation of the phosphors. There are 7 figures, 1 table and 11 Soviet references.

SURMITTED: December 17, 1959

Card 2/2

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THE REPORT OF THE PARTY OF THE

TUNITSKAYA, J.F.

USSR / Optics

Abs Jour: Referat Zhur-Fizika, 1957, No 4, 10379

Author : Levshin, V.L., Tunitskaya, V.F., Cherepneva, A.A.

Inst : Physics Institute, Academy of Sciences, USSR

Title : Origin of Localization Levels in ZnS-Cu and Co Phosphors.

Orig Pub: Optika i spektrokopiya, 1956, 1, No 2, 255-263

Abstract: An investigation was made of the thermal glow (TG) of the phosphors ZnS, ZnS-Cu, ZnS-Co and ZnS-(Cu, Co) (annealing in H_{2S}N₂ and air for 30 minutes). In ZnS, the azure glow (bands at approximately 460 millimierons) occurs only in the presence of a flux (CaCl₂). The peak of TG at -130° is due to the superstoichiometric zinc (for which favorable circumstances are produced by the chlorine), and the peak at -60° is ascribed to the caygen. The green glow is ascribed to traces of copper. In ZnS-Cu, in addition to the zinc and oxygen peaks, there appear three new peaks, barely noticeable at -5 and 0° and a considerable one at +20°. These are ascribed to copper and appear to be the cause of the longer

Card : 1/2

USSR / Optics

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Abs Jour: Referat Zhur-Fizika, 1957, No 4, 10379

afterglow of ZnS-Cu at room temperature. ZnS-Co during the instant of excitation has an azure glow, weak at room temperature, intense at -186°. Introducing Co decreases sharply the light sums from the small local levels. Simultaneously there appear deeper levels in the region of +50°. Their structure is not clear because of the small light sums. In ZnS-(Cu, Co), the peaks at -130 and -60° are suppressed rapidly with increasing Co, and the copper peaks (-5,0, and +20°) are suppressed slowly, while new peaks appear at 50 and 80°.

Card : 2/2

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GUDKOV, A.; BUSURIN, Ya.; IOFE, N.; PALKIN, G., kand. sel'khoz. nauk; TUNITSKIY, A., red.; KOROTAYEVA, D., tekhn. red.

[Manual on private livestock and poultry raising] Spravochnik po individual nomu zhivotnovodstvu i ptitsevodstvu. Koskva, Izd-vo VTsSPS Profizdat, 1946. 182 p. (MIRA 14:8) (Stock and stockbreeding) (Poultry)

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

TUNITSKIY, A.

Bryl', Yanka

"Dawn breaks in Zabolot'ye." Yanka Bryl', Reviewed by A. Tunitskiy, Mol. kolkh. 19, No. 9, 1952.

Monthly List of Russian Accessions, Library of Congress, December 1952. Unclassified.

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

TUNITSKIY, A.

White Russian Literature - History and Criticism

"Dawn breaks in Zabolot'ye." Ianka Bryl'. Reviewed by A. Tunitskiy. Mol.kolkh. 19 No. 9, 1952

Monthly List of Russian Accessions, Library of Congress, December 1952. Unclassified.

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

TUNITSKIY, L.N., kandidat fiziko-matematicheskikh nauk; IGHASHKOV, A.I., kandidat fiziko-matematicheskikh nauk

Electric discharge lighting in an elongated tube. Svetotekhnika 1 no.2:23-26 Ap '55. (NIRA 8:9)

1. Moskovskiy elektrolampovyy zavod. (Fluorescent lighting)

SKOBELEV.Y.M., inshener; TUNITSKIY_L.N., kandidat fisikomatematicheskikh nauk

Investigation of factors determining the start in fluorescent
lamps. Svetotekhnika 1 no.4:14-17 Ag '55. (MIRA 8:9)

1. Moskovskiy elektrolampovyy zavod.

(Fluorescent lamps)

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SOV/51-5-5-5/23

AU THORS:

Tatevskiy, V.M., Tunitskiy, L.M. and Novikov, M.M.

TITLE:

Vibrational Constants and Dissociation Energy of the BeF Molecule (Kolebatel'nyye postoyannyye i energiya dissotsiatsii molekuly BeF)

PERIODICAL: Optika i Spektroskopiya, 1958, Vol 5, Nr 5, pp 520-529 (USSR)

ABS TRACT:

The paper reports a new investigation of the vibrational structure of the A²II-X²I bands of BeF. The BoF spectrum was excited using gasdischarge tubes. Since the gas temperature is comparatively low in such tubes the rotational structure of the bands was weaker than in an arc and this reduced overlapping of bands and made it omsier to observe new band edges. Two types of tubes were used in this work: one working under steady-state conditions (Fig 1) and the other for pulse excitation (Fig 2). Both tubes were filled with helium under 2-3 mm Hg pressure. BeF molecules were introduced into the discharge tubes by placing some BeF2 in nickel boats inside the ubes. The tube used for steady-state discharges was U-chaped (1, in Fig 1) and had a quartz window (7, in Fig 1). The space around the electrodes (3, in Fig 1) was joined to the tube proper via liquid-hydrogen traps (4 and 5 in Fig 1). The tube was

Card 1/3

supplied from a 630 W transformer at 13 kV. The pulse-discharge source

SOV/51-5-5-5/23

Vibrational Constants and Dissociation Energy of the BeF Molecule

consisted of a straight tube (1, in Fig 2) with a fluorite window (2, in Fig 2). Its electrodes (4, in Fig 2) were supplied every second (for 0.02 sec) with a 600-700 V, 9 Å rulse. The spectra were recorded using a DFS-3 spectrograph of 2 Å/mm dispersion and 144000 resolving power. An iron spectrum was used as a wavelength standard. Measurements were made with 0.5 cm⁻¹ precision. The spectrum is shown in Fig 3. The measured band edges are given in Table 1. 48 new edges of Q1-branches and 54 new edges of R2 and R1 branches were recorded. Table 2 gives the rotational constants ω_0 , $\omega_0 x_0$ and $\omega_0 y_0$ of BeF taken from Refs 1, 3, 5, 6 and from the results reported in the present paper. Table 3 gives the values of $v_{\text{max}}^{\text{max}}$, v_{max} , v_{max} , $v_{\text{max}}^{\text{o}}$, $v_{\text{max}}^{\text{o}}$, and v_{o}^{o} , allowing for (columns I) and neglecting (columns II) the second coefficient of anharmonicity $\omega_0 y_0$ of the BeF molecule. Table 4 gives the recommended, most reliable values of the spectroscopic constants of the BeF molecule

Card 2/3

SOV/51-5-5-5/23

'Vibrational Constants and Dissociation Energy of the BeF Holecule

This table gives the dissociation energies of the $X^2\Sigma$ and $A^2\Pi$ states as 8 \pm 0.5 and 3.9 \pm 0.5 eV respectively. The author thanks L.V. Gurvich for his advice. There are 4 figures, 4 tables and 7 references,

SUBLATTED: December 6, 1957

Card 3/3

1. Beryllium fluoride--Spectrographic analysis

2. Molecules--Spectra

3. Molecules--Energy 4. Gas dishcarges--Applications

30541 8/051/60/008/06/002/024 E201/E691

5.4130 AUTHORS:

Ovcharenko, I.Ye., Tunitskiy, L.N. and Yakutin, V.I.

Analysis of the Fine Structure of the SiCl Molecular Bands TITLE:

PERIODICAL: Optika i spektroskopiya, 1960, Vol. 8, Nr 6, pp 746-751 (USSR)

ABSTRACT:

Four electron states (X2N, $B^2\Sigma$, $C^2\Delta$ and $D^2\Sigma$) of the SiCl molecule are known (Refs 1-4). The vibrational constants of these four states were reported by Jevons (Ref 4) and are listed in Table 1; the constants with question marks were considered unreliable by Jevers. The present paper reports new values of the rotational constants of the B² \sum , $X^2 \eta_{3/2}$, $X^2 \eta_{1/2}$ states, derived from the rotational analysis of the (1, 0), (0, 0) and (0, 1) bands of the $B^2\Sigma \to X^2\Pi$ transition. The spectra of SiCl were excited in a quartz pulse-discharge tube, similar to one used earlier (Ref 5) and shown schematically in a figure on p 746. The tube was filled with a mixture of silicon tetrachloride and helium. The spectra were photographed with a DFS-3 spectrograph in the third order (dispersion of 0.57 A/mm, resolving power of 432 000), and measured with a IZA-2 comparator (an iron arc spectrum was used as the wavelength standard, cf. Table 2).

Card 1/2

S/051/60/008/06/002/024 E201/E691

Analysis of the Fine Structure of the SiCl Molecular Bands

X

For the $X^2\Pi_1/2$ state the following rotational constants were obtained: $B_1 = 0.2650$, $\alpha = 0.0016$, $D_0 = 2.341 \times 10^{-7}$, $a = 0.004 \pm 0.001$ cm⁻¹; for $X^2\Pi_3/2$: $B_2 = 0.2556$, $\alpha = 0.0016$, $D_0 = 2.355 \times 10^{-7}$ cm⁻¹; for $B^2\Sigma$: B = 0.2782, $\alpha = 0.0015$, $D_0 = 1.752 \times 10^{-7}$ cm⁻¹ (Table 3). The wavenumbers of the (0.1), (0.0), (1.0) lines of the $^2\Sigma \rightarrow ^2\Pi_1/2$ transition were respectively 33662.0, 34193.6, 34892.2 cm⁻¹; for the $^2\Sigma \rightarrow ^2\Pi_3/2$ they were 33455.7, 33987.1, 34685.8 cm⁻¹, respectively. There are 1 figure, 3 tables and 6 references, of which 2 are Soviet, 3 English and 1 German.

SUBMITTED: July 20, 1959

Card 2/2

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Novikov, M.M. and Tunitskiy, L.N.

AUTHORS:

The Vibrational Constants and the Dissociation Energy of the

BeCl Molecule

PERIODICAL: Optika i spektroskopiya, 1960, Vol 8, Nr 6, pp 752-760 (USSR)

ABSTRACT:

Two electron states ($X^2\Sigma$ and $A^2\Pi$) of the BeCl molecule are known. Fredricks on and Hogan (Ref 1) determined the vibrational constants, ω_0 and $\omega_0 x_0$ and estimated the rotational constants B^*_0 and B^*_0 . Fredricks on and Hogan (Ref 1) used linear approximation to obtain the dissociation energies of the $X^2\Sigma$ (4.33 eV) and $A^2\Pi$ (3.47 eV) states. Gaydon (Ref 3) corrected the former value to 3 ± 0.5 eV. The present paper reports a new investigation of the vibrational structure of the BeCl molecular hand. A quartz pulse-discharge tube (Ref 4) was employed. The spectra (cf. a figure on p 753) were photographed with a DFS-3 spectrograph (dispersion of $2\frac{1}{4}$ /mm, resolving power of 144 000). The spectra were measured with a comparator IZA-2 (an iron are spectrum was used as the wavelength standard). Thirty new \mathbb{Q}_1 band edges and 43 new \mathbb{R}_1 and \mathbb{R}_2 band edges were recorded. Hore precise values (Table 6) of the

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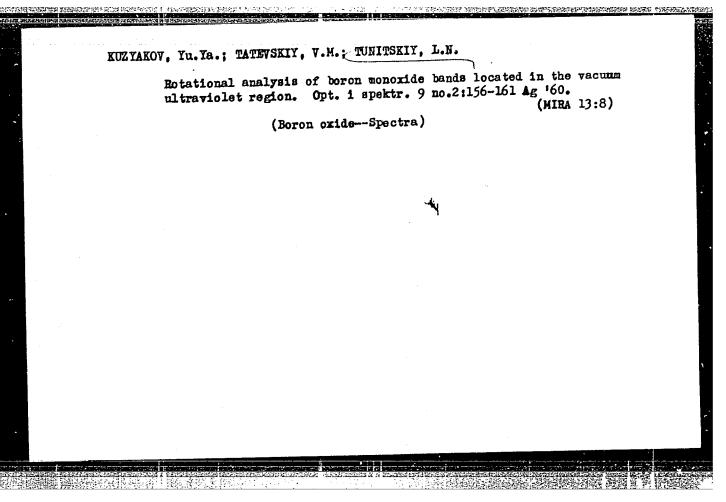
The Vibrational Constants and the Dissociation Energy of the BeCl Molecule

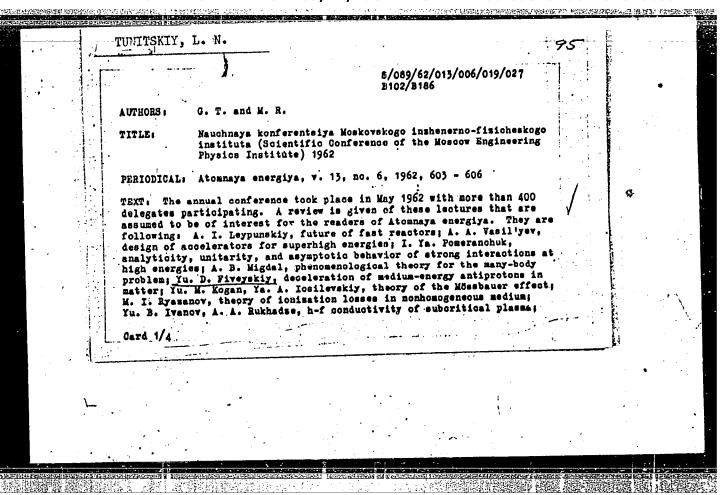
vibrational constants ω_0 and $\omega_0 x_0$ of the $X^2 \Sigma$ (841.3 and 5.11 cm⁻¹ respectively) and the $A^2\Pi$ (816.0 and 5.06 cm⁻¹ respectively) states were obtained. The second coefficients of anharmonicity $\omega_0 y_0$ were calculated for both states (0.0205 for $X^2 \Sigma$ and -0.3368 cm⁻¹ for $A^2\Pi$). The third coefficient of anharmonicity of the $X^2\Sigma$ state was found to be -0.000582 cm⁻¹. Non-linear extrapolation yielded a new and more reliable value of the dissociation energy of BeCl; 5.9 \pm 0.5 eV (Table 6). In the 2610-2620 Å region new bands were discovered (Table 7) which are due to a transition from a hitherto unknown electron state. There are 1 figure, 6 tables and 6 references, of which 2 are Soviet, 2 English, 1 French and 1 translation from English into Russian.

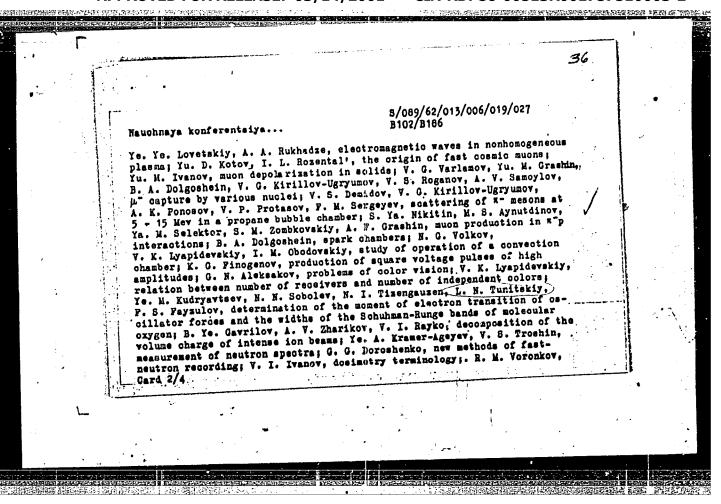
SUBMITTED: September 21, 1959

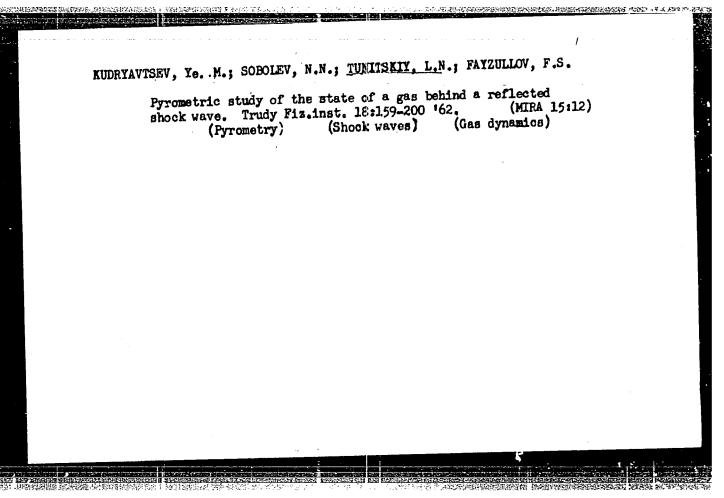
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Card 2/2.









5/051/63/014/003/005/019 E039/E120

AUTHORS:

Krindach, N.I., Kudryavisev, Ye.M., Sobolev, N.N.,

Tunitskiy, L.N., and Fayzullov, F.S.

TITLE:

Determination of the electronic transition moments of

the Schumann-Runge band system in oxygen. I. PERIODICAL: Optika i spektroskopiya, v.14, no.3, 1963, 351-361

TEXT: A method is developed for determining the square of matrix elements for electron transitions of molecules based on a measurement of the absorption in gases behind shock waves. The Schumann-Runge absorption bands for oxygen are obtained on a shock tube using helium as a working gas and equipped with apparatus for measuring temperature by the usual method of rotation of spectral lines; in this case by observing the resonance line of Ba II ($\lambda = 4554$ Å). An ionization probe is used for measuring the velocity of the front of the incident shock waves. The pulsed light source and the synchronizing circuit are described in detail. Spectra are recorded on 瓜中C-13 (DFS-13) and KC-55 (KS-55) spectrographs with glass and with quartz optics. By plotting the absorption index against wave number >> for the Card 1/2

Determination of	the electronic	S/051/63/014/00 E039/E120		
groups R(K)P(K value for the intlines of R and of the moment for this integral value. These remained in a la	- 4) and measuring tegral of the absor p form, from whi relectronic transiobtained graphical sults and the analyter paper.	thous $\begin{vmatrix} R_{\mathbf{e}}^{n} & \mathbf{m} \end{vmatrix}^2$.	The value	
There are 10 fig SUBMITTED: May	ares.			
	병원이 발발한 홍호화 사람이 되었다. 일본 사람이 보고 말았다고 있다.			
Card 2/2	맞아보다 함께 하는 것은 사실			-

KRINDACH, N.I.; SILIN-BEKCHURIN, I.A.; TUNITSKIY, L.N.; CHSRKASOV, Ye.M.

High-frequency discharge in a neon-helium laser. Ztur. teks. fil.
(MIRA 18:10)

1. Fizicheskiy institut imeni P.N.Lebedeva AN SSSR, Moskva.

L 2983-66 EWA(k)/FED/EWT(1)/EWT(m)/EPF(c)/EEC(k)-2/T/EWP(t)/EWP(k)/EWP(b)/ACCESSION NR: AP5024051 EWA(m)-2/EWA(h) SCTB/ UR/0057/65/035/009/1678 UR/0057/65/035/009/1678/1684 WP(c) 537.523.7 WG/JD AUTHOR: Krindach, N. I.; Silin-Bekchurin, I. A.; Tunitskiy, L. N.; Cherkasov Ye. 44 TITLE: Study of a high-frequency discharge in a neon-helium laser SOURCE: Zhurnal tekhnicheskoy fiziki, v. 35, no. 9, 1965, 1678-1684 gas laser, neon helium laser, hf discharge, plasma discharge TOPIC TAGS: ABSTRACT: A new method is proposed for determining the current and voltage distribution along a high-frequency discharge and a study is made of the effect on laser operation of inhomogeneities along such a discharge. The method is based on the assumption that the voltage and current at any cross section of a discharge tube can be determined by the distance of that cross section from the end of the glowing portion of discharge. This assumption holds for any stationary discharge at any cross section of which electron rise due to ionization is a unity. The experiments were carried out by means of a gas laser ($\lambda = 6328 \text{ Å}$) (see Fig. 1. of the Enclosure) which incorporated a fused-quartz discharge tube 1.7 cm long and 8 mm in diameter (internal) filled with a neon - helium mixture at a 10:1 ratio at a pressure of 0.8 mm Hg. Two plane-parallel quartz plates 01 and 0, were

L 2983-66

ACCESSION NR: AP5024051

placed at the tube ends at Brewster's angle. The equivalent circuit of the discharge tube is shown in Fig. 2. The tube was placed between the two confocal dielectric mirrors M with a 2-m radius of curvature and an - 1% transmission around 6:28 Å. The mirrors were adjusted by means of an AKT-400 collimator. The discharge tube was fed by a 30-Mc frequency from an h-f oscillator, whose voltage was supplied to 8-cm electrodes E, and E₂, while electrodes E₃, E_h, and E₅ (2.5 cm each) were grounded. The oscillator was L-coupled to the discharge tube and the currents I_1 and I2 and voltages V1 and V2 were measured by T-22 hot-wire ammeters and S-95 electrostatic voltmeters (4-pf input capacitance) respectively. The output energy was measured by means of a calibrated the mopile. The capacity of the discharge tube, varied by a movable ground rod R placed above the tube, was determined by its distance from R. In the experiments a discharge with a maximum length of 35 cm was studied. The experimental method and results are discussed in detail and indicate good agreement with computed data. Orig. art. has: 1 table and 7 figures [YK]

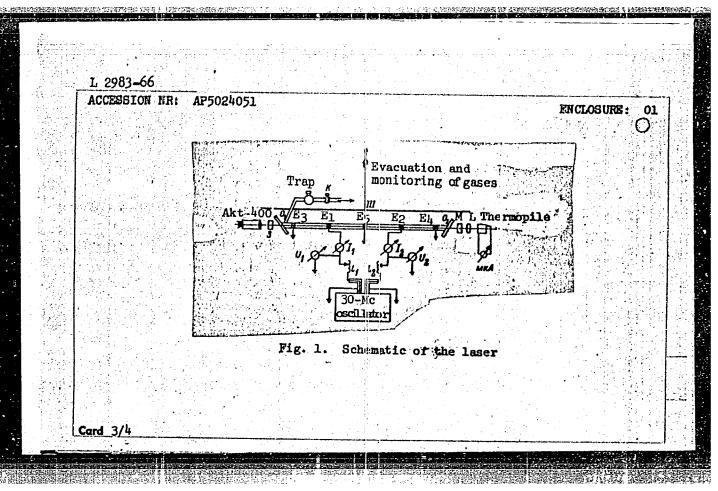
ASSOCIATION: Fizicheskiy institut imeni P. N. Lebedeva AN SSSR, Moscow (Physics

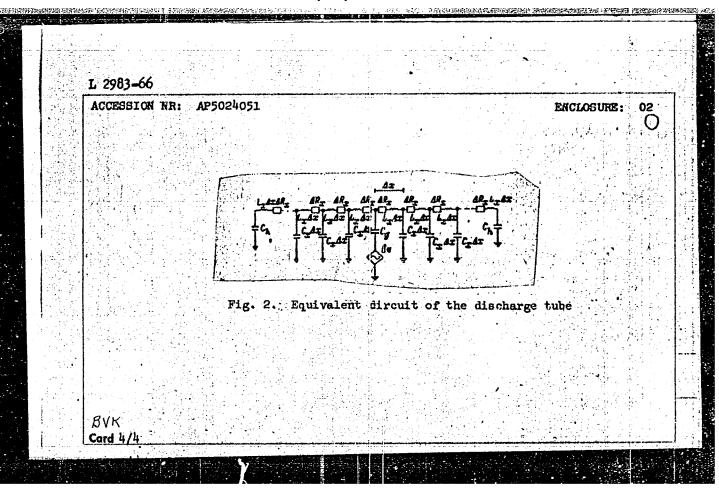
Institute, AN SSSR)

SUBMITTED: 18Jan65

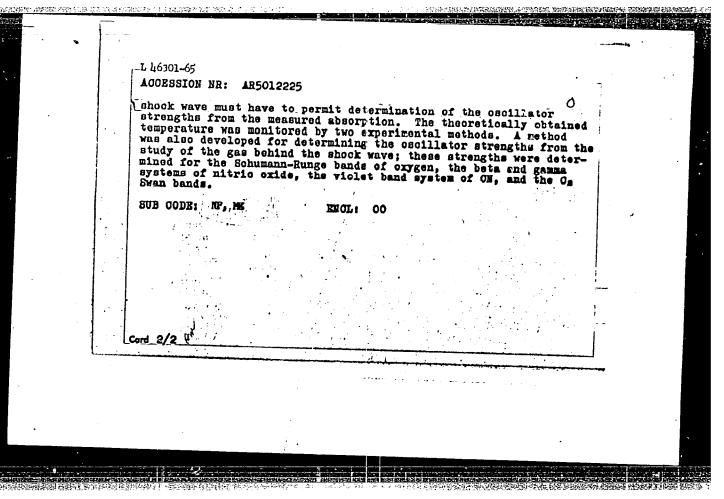
NO REF SOV: 004 ENCL: ()2 OTHER:

SUB CODE:





Card 1/2		L h6301-65 EWT(1) IJP(c) AGOESSION NR: AR5012225 SOURCE: Ref. zh. Fizika, Abs. 5D81 AUTHOR: Sobolev, N.N.; Antropov, Ve.T.; Gipoius, Ve.F. Pechenov, A.N.; Swiridov, A.G.; Unitakiy, L.N.; Fayzullov, of diatomic molecules OITED SOURCE: Tr. Komis. po spektros 64-81 TOPIC TAGS: oscillator strength, el. molecule, shook wave, oxygen, nitric TRANSLATION: To determine the oscill transitions of diatomic molecules, a coped, based on the measurement of the shook wave reflected from the end of velocity of the incident shook wave the gas behind the shook wave, it is erature and the concentration that	ectron oscillator, diatomic oxide, cyan, electronic spectration of electronic spectra experimental method was described as absorption of gas behind a fashock tube. By varying the state	trum
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ning the true values of the half-
tational structure. This method have gral absorption index which is caused a life to the control of the control

KRINDACH, N.I.; SOBOLEV, N.N.; TUNITSKIY, L.N.

Determination of the electron transition moments of the Schumann-Runge bands of the oxygen molecule. Part 3. Opt. i spektr. 15 no.5:601-608 N '63. (MIRA 16:12)

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

TUNITSKIY, L. N.

Opredeleniye momentov elektronnogo perskhoda sistem polos Shumana-Runge kisloroda report submitted for the VIIth European Congress on Molecular Spectroscopy, Budapest, 22-27 Jul 1963.

KRINDACH, N.I.; SOBOLEV, N.N.; TUNITSKIY, L.N.

Determining the electron transition moments of the Schumann-Runge bands for the oxygen molecule. Part. 2. Opt. 1 spektr. 15 no.3:298-305 S '63. (MIRA 16:10)

S/051/60/009/002/007/013/XX E201/3491

AUTHORS: Kuzyakov, Yu.Ya., Tatevskiy, V.M. and Tunitskiy, L.N.

A Rotational Analysis of Boron Monoxide Bands Lying in the Vacuum Ultraviolet Region

PERIODICAL: Optika i spektroskopiya, 1960, Vol.9, No.2, pp.156-161 Chretien (Ref.1) studied the 1300 to 2100 Å spectra of discharges in BF3 with a vacuum spectrograph fitted with a grating Chretien found several of 1 m radius and 8.3 Å/mm dispersion. bands which he ascribed to the BO molecule; the band edges, interpretations and relative intensities are given in Table 1. Later, Zelenskaya and Tunitskiy (Ref.2) reported a discharge spectrum of BF3 recorded in the vacuum ultraviolet region with a spectrograph ACC-5 (DFS-5) with a resolving power of 120000 and a dispersion of 2.7 A/mm. Zelenskaya and Tunitskiy wrongly ascribed the bands in the 1700 to 1900 Å region to a hydrogen compound of More detailed studies showed that Chretien's interpretation In the present paper the authors analyse the fine was correct. structure of the (0, 0) and (1, 0) bands (Chretien's nomenclature). A study of the isotopic shift in the BO bands, carried out in the authors' laboratory by A.A.Mal'tsev and V.M. Tatevskiy. showed that Chretien's interpretation should be corrected so that his (0, 0) and Card 1/2

s/051/60/009/002/007/013/XX

A Rotational Analysis of Boron Monoxide Bands Lying in the Vacuum

(1, 0) bands become (1, 0) and (2, 0) respectively. These bands lie between 1300 and 2100 Å and are due to 27 x25 transitions of the BO molecule; interpretation of the (1, 0) band is given in Table 2 and of the (2, 0) band in Table 3. The (Table 4)

There are here also determined (Table 4). There are 4 tables and 7 references: 2 Soviet, 3 English, 1 Swiss and 1 translation from English into Russian.

SUBMITTED: November 23, 1959

Card 2/2

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also 2308

83636 8/081/60/000/015/002/014 A006/A001

Translation from: Referativnyy zhurnal, Khimiya, 1960, No. 15, p. 15, # 60233

AUTHORS:

Kutyrkin, V.N., Peyzulayev, Sh.I., Tunitskiy, L.N.

TITLE:

Investigation of the Ber Spectrum

PERIODICAL: Fiz. sb. L'vovsk. un-t, 1957, No. 3 (8), pp. 486-489

TEXT: A # (C -3 (DFS-3) spectrograph (2A/mm dispersion) was used to investigate the BeF emission spectrum in the 2800-3400 A range (Π - Σ transition) in a discharge tube heated to 750-800°C. A reduction of the rotational structure to K \approx 15-18 as compared to the arc spectrum (K \gg 55), made it possible to obtain a considerably greater number of band edges than in operating with an arc. Wave numbers of 1-1 band lines are given. The presence of lines with K = 0 and 1 in the branch show that the BeF molecule terms are inverted ones. See also RZhKhim, 1959, No. 8, # 26114.

A. Mal'tsev

Translator's note: This is the full translation of the original Russian abstract.

Card 1/1

83637

5.2400A

S/081/60/000/015/003/014 A006/A001

Translation from: Referativnyy zhurnal, Khimiya, 1960, No. 15, p. 15, # 60234

AUTHORS: Zelenskaya, L.G., Tunitskiy, L.N.

TITLE: Investigation of a Discharge Spectrum in BF3 to Vacuum Ultraviolet

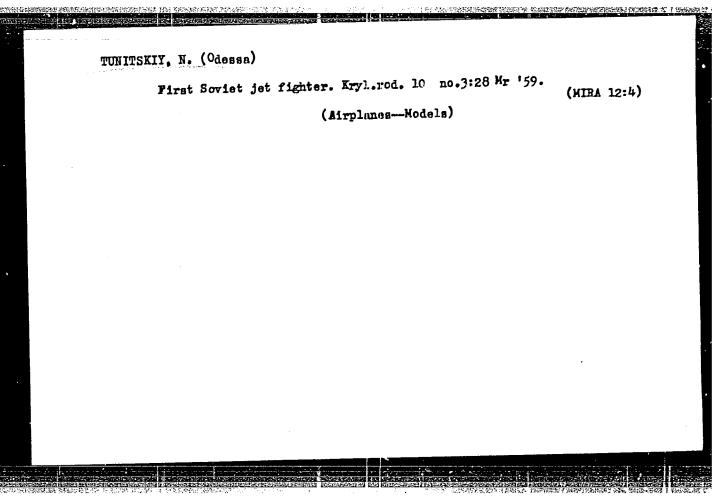
PERIODICAL: Fiz. sb. L'vovsk, un-t, 1957, No. 3 (8), pp. 489-493

TEXT: A Π ϕ (-5 (DFS-5) vacuum spectrograph (2.7 A/mm dispersion) was used to investigate the discharge emission spectrum in BF3. The rotational structure of two bands in the 1700-1900A range was determined. The bands must according to Cretenin's data (Cretenin, Helv. phys. acta, 1950, Vol. 23, p. 259) belong to the $^{2}\Pi$ - $^{2}\Sigma$ transition of the BO molecule, appearing as a contamination. As a result of rotational analysis it is shown that it is probably wrong to relate these bands to the BO molecule, and they are related to the $^{1}\Pi$ - $^{1}\Pi$ or $^{2}\Sigma$ - $^{2}\Sigma$ transition of some hydrogen compound. Rotational constants B' and B' \approx 8.7 cm⁻¹A are determined.

A. Mal'tsev

Translator's note: This is the full translation of the original Russian abstract.

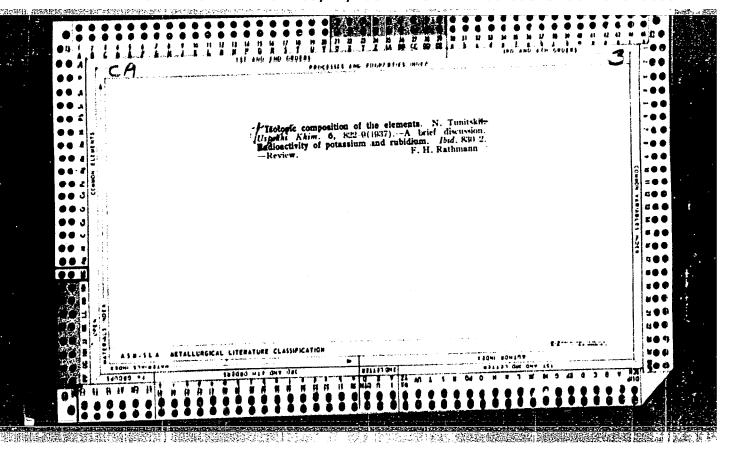
Card 1/1

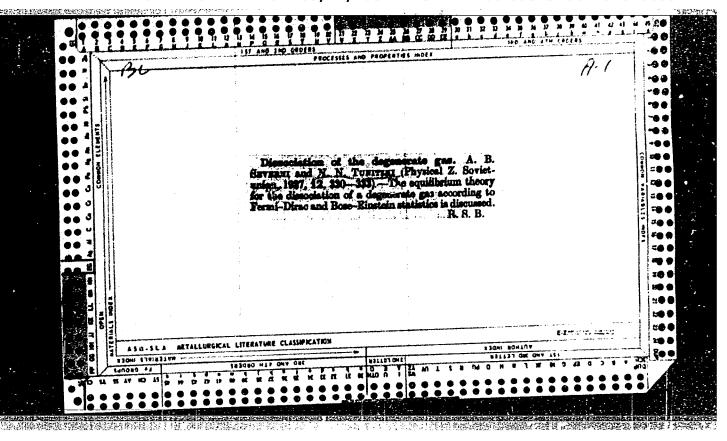


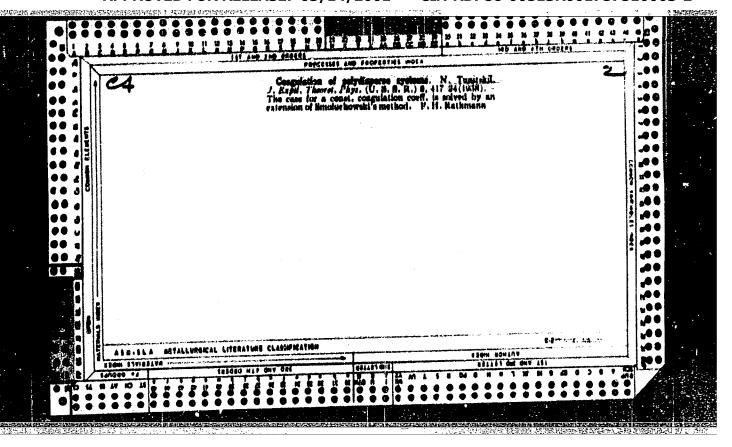
SUMIN, I..V.; GUE'YEV, M.V.; TUNITSKIY, N.N.

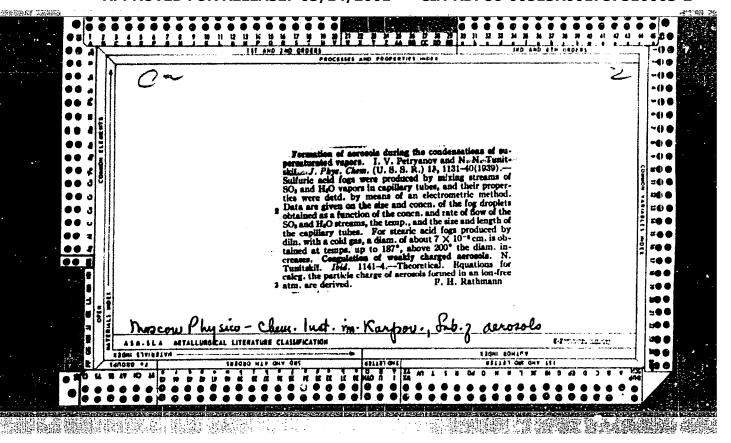
Average time of formation of fragment ions from n-hezane. Kin.: kat. 5 no.6:961-967 N-D *64. (MIRA 18:3)

1. Nauchno-issledovatel'skiy Fiziko-khimicheskiy institut imeni Karpova, Moskva.









TUNITSKIY4N8

600

1. TUNITSKIY, N.

2. USSR (600)

"The Coagulation of Weakly (slabo) Charged Aerosols", Zhur. Fiz. Khim, 13, No. 8, 1939. Moscow, Physico-Chemical Institute imeni L. Ya. Karpov, Lab. of Aerosols. Received 27 Feb 1939.

9. Report U-1615, 3 Jan 1952.

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

BYKOV, R.I.; MAL'TSEVA, A.K.; TURANOV, V.A.; GAVRILOV, V.P.

Regularities in the distribution of oil and gas fields in the
Jurassic sediments of the central part of the Turan Plateau.

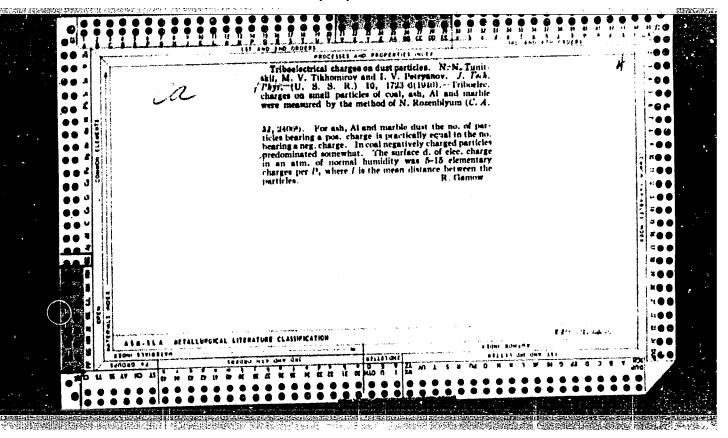
Trudy MINKHIGP no.43:125-134 '63. (MIRA 17:4)

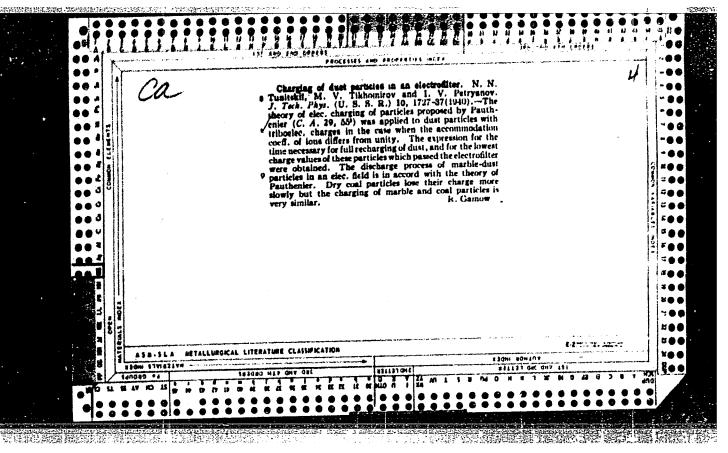
TURBIN, N. V.; MIRONENKO, A. V.

*The genetic blocking of alkaloid biosynthesis in varieties of forier lupin." report submitted for 10th Intl Botanical Cong, Edinburgn, 3-12 Aug 64.

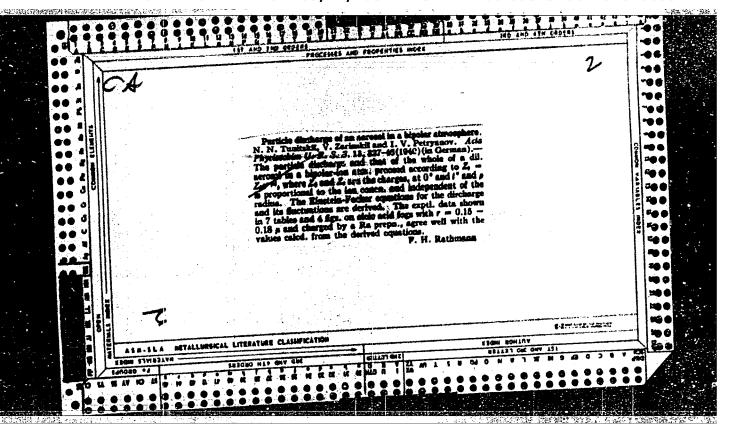
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APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"





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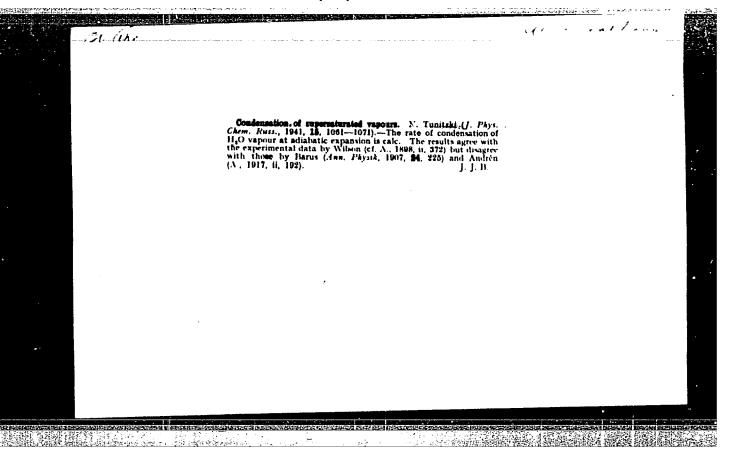
TUNITSKIY, N.; ZARINSKIY, V.; PETRIALLY, I.

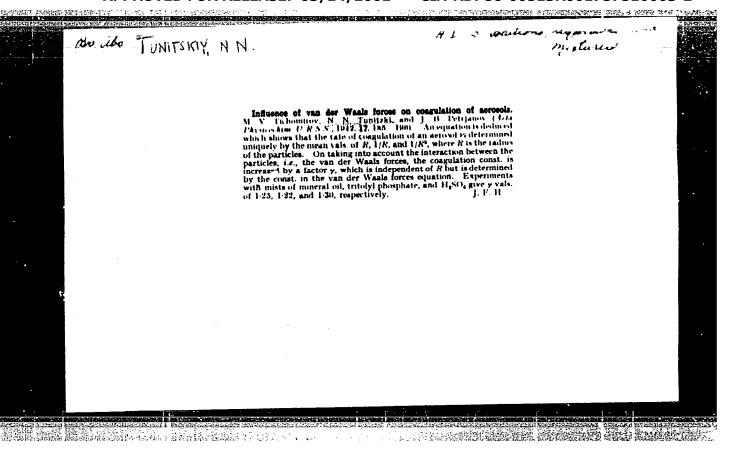
Physico-Chemical Institute imeni L. Ya. Karpov, (-1940-).

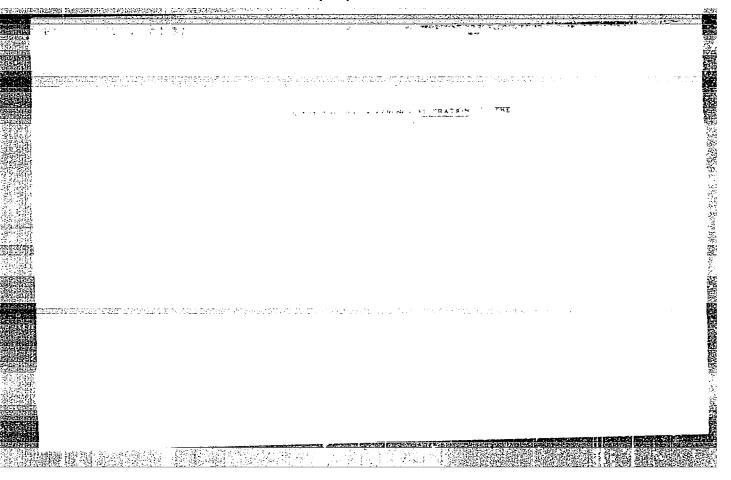
"The Charging of Aerosols in a Bipole-Ionized Atmosphere."

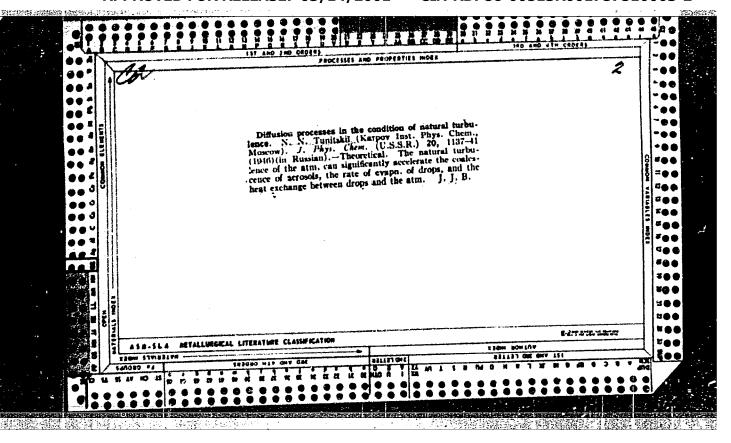
Zhur. Fiz. Khim., Vol. 14, No. 4, 1940.

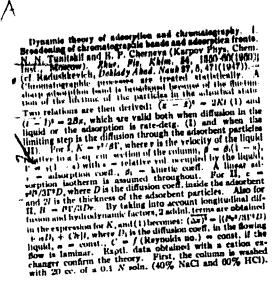
APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"







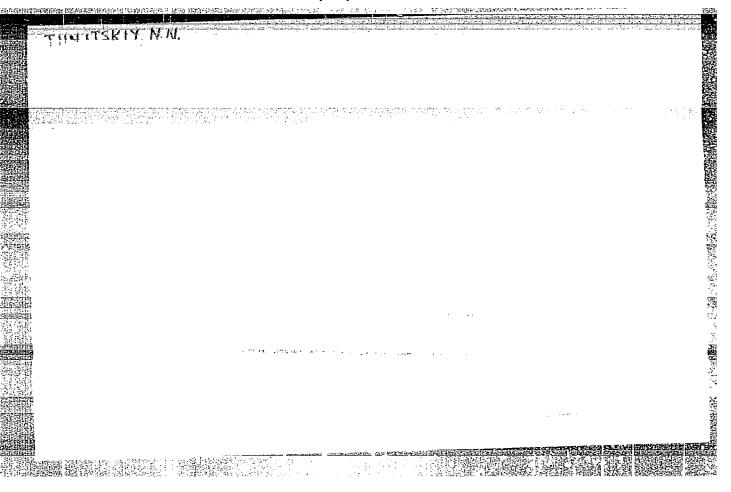




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then the band is displaced with a 0.1 N Bi Lache pointed at a given rate, and samples (12 cc.) are taken and analyse (14 ct.) of NaCl). The above formulas give $(\Delta V)^2/V = C_0(2)$ But $(\Delta V)^2/V = ((\overline{\Delta N})^2/\overline{N})\omega$ with $\overline{N} = 2N_{CN}/2C_N$ and $(\overline{\Delta N})^2 = [2(N-N)^2C_N/2C_N]$. N= no. of sample, $\omega=$ its vol., $C_N=$ its NaCl conen. A straight line is obtained in a diagram $(\overline{\Delta V})^2/V$ against v in the range 0.3 < v < 2n, inin., as predicted by (2). The theory also predicte that the distribution function W(u,t) will be Gaussian at least for sufficiently large values of t. This is verified by plotting $[\log(100/\epsilon]^{1/\epsilon}$ against $\Delta N=N-N$; a straight line is obtained, as expected for a Gaussian distribution.

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	Tunitskii ai	Doklady Akad, Naus 3.3.3.	ographic strip			
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KOLOTYRKIN, V.M.; TIKHOMIROV, M.V.; TUNITSKIY, N.M.; SEMENOV, N.N., akademik.

Mass spectrum of methane at increased pressure. Dokl.AN SSSR 92 no.6:1193-1195 0 '53. (MLRA 6:10)

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1. Akademiya nauk SSSR (for Semenov). 2. Fiziko-khimicheskiy institut im. L.Ya.Karpova (for Kolotyrkin, Tikhomirov and Tunitskiy). (Methane) (Spectrum analysis)

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"

Tunitsky, N.N.

USSR/ Physics - Physical chemistry

Card 1/1 : Pub. 147 - 18/22

Authors : Tunitskiy, N. N.; Cherneva, E. P.; and Andreyev, V. I.

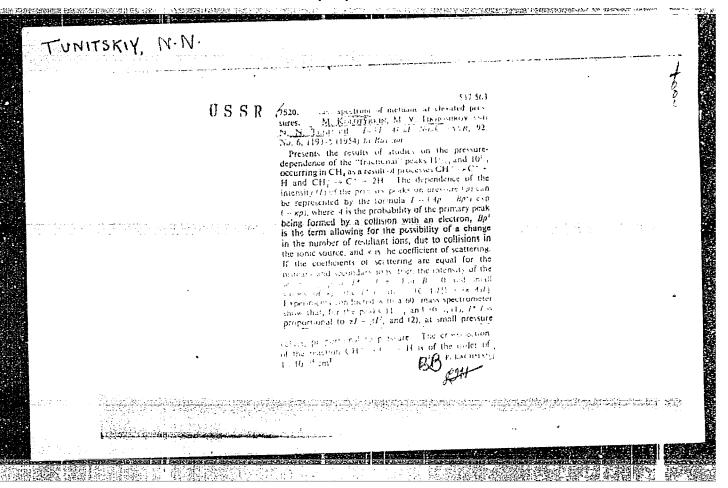
Title : On the theory of the dynamics of sorption and chromatography. Part 3.—
The dynamics of ion-exchange sorption during intradiffusion kinetics.

Periodical: Zhur. fiz. khim. 28/11, 2006-2020, November 1954

The dynamics of ion-exchange sorption was investigated to determine the behavior of the sorbent layer in a flow containing the sorption substance. The conditions favorable for parallel transfer for ion-exchange sorption are indicated. The term expressing the time loss of the protective effect is introduced. The role of internal diffusion and hydrodynamic factors in the sorption dynamics, is exclaimed. The experimental data obtained for calcium sorption with cationite in H-form, hydrogen sorption with cathonite in Ca-form and sorption of radioactive calcium with cationite in Ca-form were found to be in excellent conformity with the theoretical data. The live USSR references (1929-1953). Tables; graphs.

Institution: The L. Ya. Karpov Physico-Chemical Institute, Noscou

Submitted :- Narch 26, 195/4



LUNITSKIY,

USSR/Chemistry - Physical chemistry

Gard 1/1

Pub. 22 - 24/45

Authors

: Tunitskiy, N. N.

: The layer method of calculating chromatographic processes

Periodical : Dok. AN SSSR 99/4, 577-579, Dec 1, 1954

Abstract

: The layer method for the calculation of chromatographic separation processes, based on the assumption that the continuous process in the column can be considered as a series of equilibriums in the small layers of a sorbent, is described. Some results obtained from calculations in accordance with the layer method are listed. Seven references: 6-USSR and 1-USA (1947-1953).

Institution: The L. Ya. Karpov Physico-Chemical Institute

Presented by: Academician M. M. Dubinin, July 3, 1954

"APPROVED FOR RELEASE: 03/14/2001

CIA-RDP86-00513R001757510003-2

JunitsKix, N. A

USSR/Physical Chemistry - Photochemistry. Radiation Chemistry. Theory of the Photographic Process, B-10

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 470

Tunitskiy, N. N., Kupriyanov, S. Ye., and Tikhomirov, M. V. Author:

Institution: Academy of Sciences USSR

Title: Effect of Electrons of Different Energies on the Ionization and Dissociation of Some Hydrocarbon Halides

Original

Periodical: Sb. rabot po radiatsionnoy khimii, Moscow, Publishing House of the

Academy of Sciences USSR, 1955, 223-240

The ionization and dissociation of halogen derivatives of hydrocar-Abstract:

bons has been carried out with a type MS-1 mass spectrometer, supplemented as follows: (1) automatic scanning of the mass spectrum, (2) automatic recording of the mass spectrum, and (3) introduction of the sample into the ion source. The mass spectra of CH4, CH3Cl, CH2Cl2, CHCl3, CCl4, CH3Br, and CH3I have been recorded with electron energies of 100 ev. It is shown that as the number of helide atoms

Card 1/2

USSR/Physical Chemistry - Photochemistry. Radiation Chemistry. Theory of the Photographic Process, B-10

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 470

ment ions.

Abstract: in the molecule increases, the maximum intensity of the ionic current shifts from the region of molecular ions to that of dons from by the splitting off of an atom, i.e., dissociation begins to overshadow ionization. Anomalous ions and some secondary processes which occur during ionization and dissociation were investigated. The formation of H2X† was observed in the mass spectra of some halogen derivatives of methane. The dependence of the ionization and dissociation of the molecules CH3I, C2H2Cl4, CH3Br, and CH3Cl on the electron energy (up to 1,100 ev) was investigated. It is shown that as the energy of the ionizing electrons increases, the mass spectra contained fewer frag.

Card 2/2

KUPRIYANOV, S.Ye.; DZHAGATSPANYAN, R.V.; TIKHOMIROV, M.V.; TUNITSKIY, E.N.

Mass-spectrometric method of analysing chlorins derivatives of
methane. Zav.lab.21 no.l0:1182-1188 '55. (MIRA 9:1)

1. Mauchno-issledovatel'skiy fiziko-khimicheskiy institut imeni
L.Ya.Karpova.

(Mass spectrometry) (Methane)

TUNITSKIY, NY

USSE/Physics - Physical chemistry

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Card 1/2

Rub. 22 - 32/51

Authors

Tikhomirov, M. V.; Kolotyrkin, V. M.; and Tunitskiy, N. H.

Title

About the dissociation of primary ions in a mass-spectrometer

Periodical :

Dok. AN SSSR 101/5, 903-905, Apr 11, 1955

Abstract

The relation between the intensity of "fractional" n-butane peaks and pressure was investigated to explain the mechanism of primary ion dissociation at greater pressures. It is pointed cut that the dissociation at greater pressures. It is pointed cut that the dissociation during collision, as in the case of spontaneous decomposition, may depend upon the ion excitation and that the excitation varies departing upon the energy of the ionizing electrons. It was found that the relative intensity of the "fractional" peaks increases with the electron

Institution : Presented by:

The A. A. Zhdanov State University, Leningrad Academician A. N. Terenin, November 14, 1954

Card 2/2 Pub. 22 - 32/51

Periodical: Dok. AN SSSR 101/5, 903-905, Apr 11, 1955

Abstract: energy, this is due to the fact that the spontaneous decomposition of the ions and their decomposition depend in various degrees upon the electron energy. Eight references: 3 German, 2 UESR, 2 USA and 1 English (1939-1953). Graphs.

USSE/Physics - Physical chemistry

Card 1/1

Pub. 22 - 28/47

Authors

Tunitskiy, N. N.; Smirnova, R. M.; and Tikhomirov, M. V.

Title

About "broken" peaks in the mass spectrum of hydrogen

Periodical

Dok. AN SSSR 101/6, 1083 - 1084, Apr. 21, 1955

Abstract

* A 60° mass spectrometer of low resolving power and magnetic sweep of the mass spectrum was employed in determining the relation between the cross sections of H₂⁺ ion dissociation and the energies of the ionizing electrometer. It is shown that the dissociation process results in the formation of a blurred peak (band) in the mass spectrum of hydrogen at an apparent mass of 1/2. It was established that the (1/2) peak, corresponding to dissociation protons, has a width approximately 5 times greater than the basic peak. The relation between dissociation cross section and ion energy is explained. Five references: 4 USSR and 1 German (1939-1954). Graphs.

Institution: The L. Ya. Karpov Sc. Res. Phys. Chem. Inst.

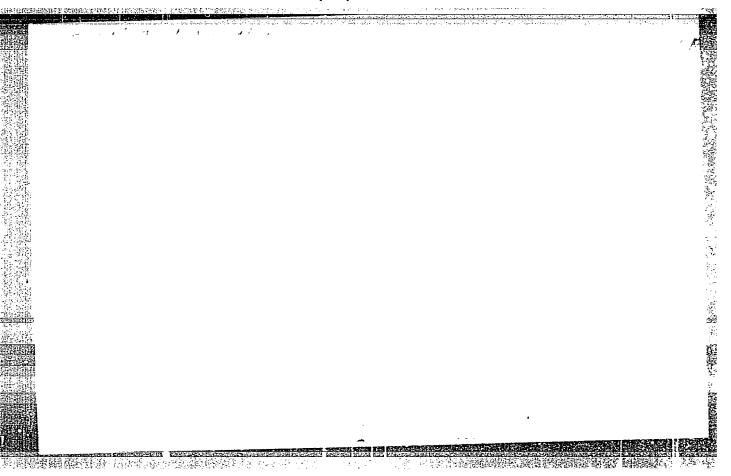
Presented by: Academician V. N. Kondratyev, November 12, 1954

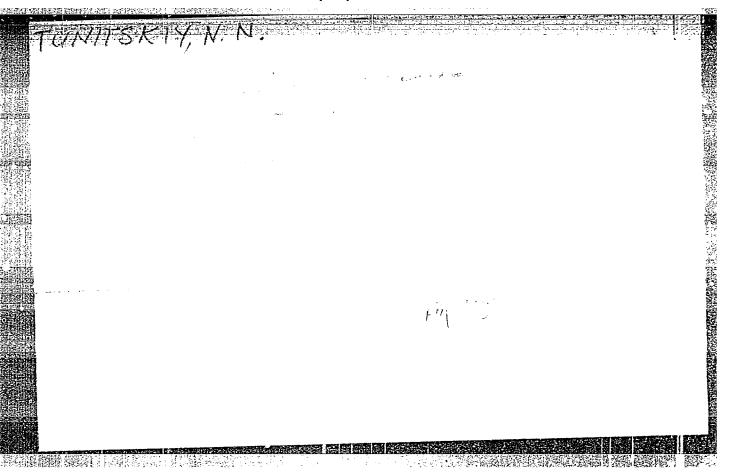
DZHAGATSPANYAN, R.V.; TUNITSKIY, N.N. THE PARTY NAMED IN COLUMN Experimental study of the dilution of chromategraphic bands.

Dokl.AN SSSR 105 no.6:1282-1284 D 155. (HIRA 9:4)

1.Fizike-khimicheskiy institut imeni L.Ya.Karpeva. Predstavlene akademikem V.A.Karginym. (Chremategraphic analysis)

APPROVED FOR RELEASE: 03/14/2001 CIA-RDP86-00513R001757510003-2"





TUNITSKIY, N. N. (Prof.)

"The Molecule- and Ionic Dissociation in the Mass Spectrometer."

report presented at Scientific Conference at the Inst. for Ph sical Chemistry imeni L. Ya. Karpov, Acad. Sci. USSR, Nov 1957.

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SOV/137-58-7-14194

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 7, p 33 (USSR)

AUTHOR: Tunitskiy, N. N.

TITLE: Theory of Ion-exchange Processes (Teoriya ionoobmennykh

protsessov)

PERIODICAL: V sb.: Materialy Soveshchaniya po primeneniyu ionnogo

obmena v tsvetn. metallurgii. Moscow, 1957, pp 4-16

ABSTRACT: Bibliographic entry

1. Ion exchange--Theory

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SOV/137-58-9-18449

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 9, p 38 (USSR)

AUTHORS: Cherneva, Ye. P., Tunitskiy, N. N.

TITLE: Diffusion Through Ion-exchange Membranes (Diffuziya cherez

ionnoobmennyye membrany)

PERIODICAL: V sb. Materialy Soveshchaniya po primeneniyu ionnogo obmena v tsvetn. metallurgii, Moscow, 1957, pp 43-47

obmena v isvein. metaliargi, meson, iii

ABSTRACT: The ranges of technical application of ion-exchange membranes

are traced. The rates of permeation of cations through a cationite membrane on both sides of which different solutions

were present initially were studied.

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1. Ion exchange--Theory

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事態體調整 國海道特別自己公司行為自己

TUNITSKIY, N.N.; TIKHOMIROV, M.V.; KUPRIYANOV, S.Ye.; KOLOTYRKIN, V.M.;
GUR'YEV, M.V.; POTAPOV, V.K.

Studies in the field of mass spectrometry. Probl.fiz.khim. no.1:122-128 '58. (MIRA 15:11)

l. Laboratoriya adsorbtsionnykh protsessov Nauchnoissledovatel'skogo fiziko-khimicheskogo instituta im. Karpova.

(Mass spectrometry)

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TUNITSKIY, N.N.

21(8)

PHASE I BOOK EXPLOITATION

sov/1140

- * Resoyuznoye soveshchaniye po radiatsionnoy khimii. lst, Moscow, 1957.
 - Trudy (Transactions of the First Conference on Radiation Chemistry) Moscow, Izd-vo AN SSSR, 1958. 330 p. 4,000 copies printed.
 - Sponsoring Agencies: Akademiya nauk SSSR. Otdeleniye khimicheskikh nauk, and U.S.S.R. Ministerstvo khimicheskoy promyshlennosti.
 - Editorial Board: Bakh, N.A. Professor (resp. ed.); Medvedev, S.S., Corresponding Member, Academy of Sciences, USSR, Veselovskiy, V.I., Professor, Dolin, P.I., Doctor of Chemical Sciences; Miller, N.B., Candidate of Chemical Sciences, Tsetlin, B.L., Candidate of Chemical Sciences (Secretary). Eds. of Publishing House: Trifonov, D.N. and Bugayenko, L.T.; Tech. Ed.: Moskvicheva, N.I.
 - PURPOSE: This book publishes the reports of the First All-Union Conference on Radiation Chemistry in Moscow, March 25 - 30.
 - COVERAGE: This collection includes fifty-seven reports and follow up discussions of each sub-group of reports classified as follows:

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Transactions of the First (Cont.)

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- primary functions in radiation-chemical processes,
- radiation chemistry of water solutions (inorganic and organic systems),

radiation-electrochemical processes,

- the effect of radiation on substances which take part in biochemical processes,
- radiation chemistry of simple organic systems,

radiation effects on polymers, and

sources of radiation.

According to the editors, the discussions reveal differences in points of view of Soviet scientists on various problems of radiation chemistry; specifically, the mechanism of the action of radiation on concentrated water solutions, the practical importance of radiation-galvanic phenomena, the mechanism of the action of radiation on polymers, etc. The editors also the action of radiation on polymers, etc. note that the conference revealed inadequate development in some important areas of radiation chemistry, Particularly the theory of initiation of radiolysis, and the action of radiation on solid bodies.

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